In the Claims:

Listing of the Claims:

Please amend claims 1 and 26 as indicated. Please add claims 32 and 33 as indicated.

- (Currently Amended) A vaccine useful for the treatment of melanoma comprising
 irradiated autologous melanoma cells conjugated to a hapten, said hapten selected from
 the group consisting of trynitrophenyl trinitrophenyl and N-iodoacetyl-N'-5 sulfonic 1naphtyl ethylene diamine; and mixed with an immunological adjuvant, wherein said
 immunological adjuvant is Bacille Calmette-Guerin.
- 2. (Previously Presented) A method for treating melanoma comprising administering cyclophosphamide followed by intradermal administration of a therapeutically effective amount of the vaccine of claim 1, wherein said vaccine induces a delayed-type hypersensitivity (DTH) response against unmodified melanoma cells.
- 3. Canceled
- 4. Canceled
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- 20. Canceled

- 21. (Previously Presented) The vaccine of claim 1, wherein said autologous melanoma cells are cryopreserved.
- 22. (Previously Presented) The method of claim 2, wherein said autologous melanoma cells are cryopreserved.
- 23. (Previously Presented) The method of claim 2, wherein said vaccine is injected into three contiguous sites on an upper arm or leg.
- 24. (Previously Presented) The method of claim 2, wherein said vaccine is administered to post-surgical melanoma patients.
- 25. (Previously Presented) The method of claim 2, wherein said vaccine is administered to stage four melanoma patients.
- 26. (Currently Amended) A method for treating melanoma comprising administering cyclophosphamide followed by intradermal administration of a therapeutically effective amount of a vaccine composition comprising autologous irradiated melanoma cells conjugated to a hapten, said hapten selected from the group consisting of dinitrophenyl, trinitrophenyl, and N-iodoacetyl-N'-5 sulfonic 1-naphtyl ethylene diamine and mixed with Bacille Calmette-Guerin, wherein administration of said vaccine induces a delayed-type hypersensitivity (DTH) response against unmodified melanoma cells.
- 27. (Previously Presented) The method of claim 26, wherein said vaccine is injected into three contiguous sites on an upper arm or leg.
- 28. (Previously Presented) The method of claim 26, wherein said vaccine is administered to post-surgical melanoma patients.
- 29. (Previously Presented) The method of claim 26, wherein said vaccine is administered to stage four melanoma patients.
- 30. (Previously Presented) The method of claim 26, wherein said vaccine is administered every four weeks.
- 31. (Previously Presented) The method of claim 26, wherein said autologous melanoma cells are cryopreserved.
- 32. (New) The vaccine of claim 1, further comprising an immunomodulating drug.
- 33. (New) The vaccine of claim 32, wherein the immunomodulating drug is IL-2.